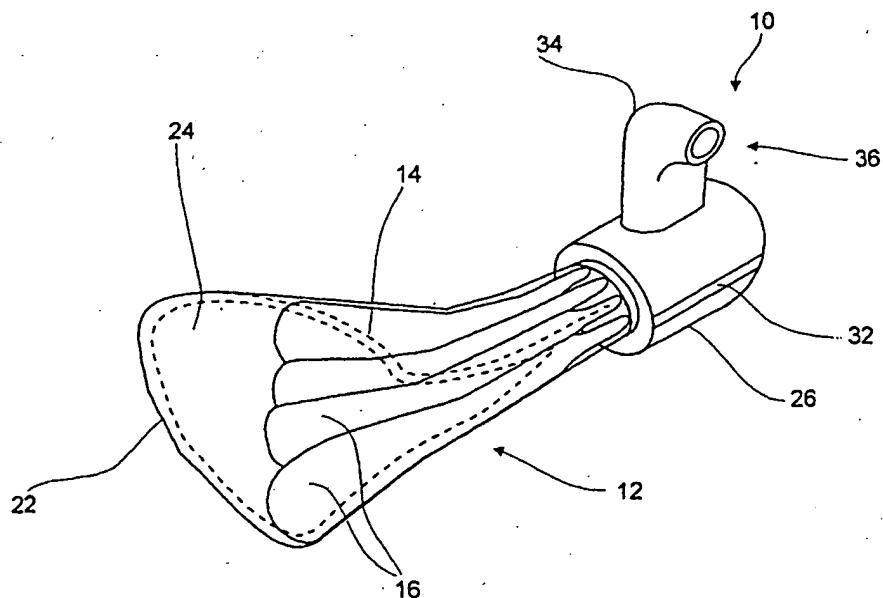




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(54) Title: MINIMAL INVASIVE CARDIAC MASSAGE DEVICE



## (57) Abstract

A cardiac massage device (10) deployable by a minimal invasive procedure for applying pulsatory pressure to a ventricular wall of a heart includes an inflatable receptacle (12) deployable between a closed insertion state and an open state. The open state of the device is configured for substantially enveloping the ventricular portion of the heart. A plurality of support elements (14) are mechanically linked to the inflatable receptacle so as to deploy the inflatable receptacle to its open state. Preferably, the support elements are implemented as flexible rods resiliently biased to return to a pre-defined shape and a retaining mechanism (22) is provided. Also described is a deployment technique employing suction clamping and a two-stage trocar (40, 42).

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## Minimal Invasive Cardiac Massage Device

### FIELD AND BACKGROUND OF THE INVENTION

The present invention relates to devices for cardiac treatment, and more particularly to a device for cardiac massage which can be positioned by a minimal invasive procedure.

Various devices are known for cardiac massage. Examples are disclosed in U.S. Patents Nos. 4,536,893 to Parravicini and 5,713,954 to Rosenberg et al. These describe inflatable cup-shaped devices which are deployed around the heart. A pulsating supply of fluid to the devices generates inward pressure on the ventricular portion of the heart, thereby assisting the pumping action.

These devices are both designed for implantation during an open heart surgery procedure. As a result, their use is limited to long term applications in which the patient is in a sufficiently fit general state of health to be able to withstand such a procedure. Furthermore, such implantable devices do not allow sufficiently rapid deployment to be useful in cases for which cardiac assist must be started very rapidly or for emergency applications such as Cardio-Pulmonary Resuscitation (CPR) procedures where direct cardiac massage could be much more effective than the conventional procedure of pressing on the rib cage.

A further shortcoming of the existing cardiac massage devices relates to the manner of retaining the devices in position around the heart. It is important that the device applies pressure only to the ventricular region of the heart. However, when pressure is applied from a device surrounding only the lower part of the heart

corresponding to the ventricular region, the shape and texture of the heart tends to make it slip out of the device. To counter this tendency, the device of Parravicini provides straps for wrapping around the root of the aorta and pulmonary artery. This somewhat primitive fastening technique is, however, clearly limited to open 5 surgical procedures and may exert pressure on the aorta or atria.

There is therefore a need for a device for cardiac massage which may be deployed rapidly in a minimal invasive procedure and which provides effective retention of the device in position around the heart.

#### SUMMARY OF THE INVENTION

10 The present invention is a device for cardiac massage which can be positioned by a minimal invasive procedure.

According to the teachings of the present invention there is provided, a cardiac massage device deployable by a minimal invasive procedure for applying pulsatory pressure to a ventricular wall of a heart, the device comprising: (a) an 15 inflatable receptacle deployable between a closed insertion state and an open state, the open state being configured for substantially enveloping the ventricular portion of the heart; and (b) a plurality of support elements mechanically linked to the inflatable receptacle so as to deploy the inflatable receptacle to the open state.

According to a further feature of the present invention, the support elements 20 are implemented as flexible rods resiliently biased to return to a pre-defined shape.

According to a further feature of the present invention, there is also provided an elongated hollow structure for storage and deployment of the device,

the inflatable receptacle and the flexible rods being retractable within the elongated hollow structure so as to confine the flexible rods to an initial state corresponding to the closed insertion state of the inflatable receptacle, the flexible rods returning to the pre-defined shape as they advance out from the elongated hollow structure.

5 According to a further feature of the present invention, the elongated hollow structure forms a storage stage of a two-stage trocar, an insertion stage of the trocar being configured for insertion through the skin of a patient as part of a minimal invasive surgical procedure.

According to a further feature of the present invention, the insertion stage of  
10 the trocar includes a curved hollow guide portion for insertion within the patient, the curved hollow guide portion providing a turn angle of between about 15° and about 30°.

According to a further feature of the present invention, there is also provided a manually operable mechanism for displacing the inflatable receptacle  
15 and the flexible rods along a length of the trocar.

According to a further feature of the present invention, there is also provided a retaining mechanism associated with the inflatable receptacle and configured for deploying around a supraventricular portion of the heart so as to retain the inflatable receptacle in a position enveloping the ventricular portion of  
20 the heart.

According to a further feature of the present invention, the retaining mechanism includes at least one drawstring mechanism.

According to a further feature of the present invention, the at least one drawstring mechanism is connected to the inflatable receptacle via a non-inflatable web.

According to an alternative feature of the present invention, the retaining 5 mechanism includes at least one inflatable retaining element, the inflatable retaining element and the inflatable receptacle being provided with independent fluid supply conduits.

According to a further feature of the present invention, there is also provided a retractable suction clamping system including: (a) a suction cup 10 deployed for suction clamping to the apex of the heart; and (b) an axially displaceable tube forming a fluid connection with the suction cup, such that, when the tube is displaced forward and suction is applied to the tube, the suction cup clamps to the heart and, when the suction is released and the tube is displaced rearwards, the suction cup is withdrawn from contact with the heart.

15 According to a further feature of the present invention, there is also provided at least one electrode deployed on an inner surface of the inflatable receptacle so as to contact the heart when the device is deployed.

There is also provided according to the teachings of the present invention, a 20 cardiac massage device deployable by a minimal invasive procedure for applying pulsatory pressure to a ventricular wall of a heart, the device comprising: (a) an inflatable receptacle for substantially enveloping the ventricular portion of the heart, the inflatable receptacle being configured to exert pulsating pressure against the ventricular wall in response to a pulsating fluid supply; and (b) a retaining

mechanism associated with the inflatable receptacle and configured for deploying around a supraventricular portion of the heart so as to retain the inflatable receptacle in a position substantially enveloping the ventricular portion of the heart.

5 According to a further feature of the present invention, the retaining mechanism includes at least one drawstring mechanism.

According to a further feature of the present invention, the at least one drawstring mechanism is connected to the inflatable receptacle via a non-inflatable web.

10 According to an alternative feature of the present invention, the retaining mechanism includes at least one inflatable retaining element, the inflatable retaining element and the inflatable receptacle being provided with independent fluid supply conduits.

There is also provided according to the teachings of the present invention, a  
15 method of minimal invasive deployment of a cardiac massage device around the ventricular portion of the heart of a patient, the method comprising the steps of:  
(a) introducing the device in a compact state through the pericardium in the region near the diaphragm; and (b) opening the device to form a receptacle which substantially envelopes the ventricular portion of the heart.

20 According to a further feature of the present invention, the opening of the device occurs through a plurality of resiliently biased flexible rods deployed within the device returning to a pre-defined shape.

According to a further feature of the present invention, suction clamping is also applied to the surface of the heart during deployment of the device, the suction clamping being removed once the device is properly deployed.

According to a further feature of the present invention, a retaining mechanism is deployed around a supraventricular portion of the heart so as to 5 retain the receptacle in a position enveloping the ventricular portion of the heart.

According to a further feature of the present invention, the compact state exhibits a maximum diameter of no more than about 4 cm and preferably between about 1.5 and about 2.5 cm.

#### 10 BRIEF DESCRIPTION OF THE DRAWINGS

The invention is herein described, by way of example only, with reference to the accompanying drawings, wherein:

FIG. 1 is a schematic isometric view of a device for cardiac massage, constructed and operative according to the teachings of the present invention, ready 15 to be deployed;

FIG. 2 is a schematic isometric view of the device of Figure 1 in a deployed state;

FIG. 3 is a schematic cross-sectional view through the deployed device of Figure 2;

20 FIG. 4 is a schematic cross-sectional view through a base connector of the device of Figure 2;

FIG. 5 is a schematic isometric view of the device of Figure 2 engaged around a heart;

FIG. 6 is a schematic isometric view of a two-stage trocar for use in deploying the device of Figure 1;

5 FIG. 7 is a longitudinal cross-sectional view through the trocar of Figure 6 with the device of Figure 1 stored therein;

FIG. 8 is a schematic isometric view of the first stage of the trocar of Figure 6 with the device of Figure 1 in its deployed state;

10 FIGS. 9A-9D are schematic side views showing a sequence of stages during blind deployment of the device of Figure 1 around a heart;

FIG. 10 is a schematic rear view of a first implementation of a retaining mechanism for the device of Figure 1 employing a drawstring mechanism; and

15 FIG. 11 is a schematic isometric view of a second implementation of a retaining mechanism for the device of Figure 1 employing an inflatable retaining element.

#### DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is a device for cardiac massage which can be positioned by a minimal invasive procedure.

20 The principles and operation of a device according to the present invention may be better understood with reference to the drawings and the accompanying description.

Referring now to the drawings, Figures 1-5 show a cardiac massage device, generally designated 10, constructed and operative according to the teachings of the present invention, which is deployable by a minimal invasive procedure. Device 10 is used to apply pulsatory pressure to a ventricular wall of a heart.

5        Generally speaking, cardiac massage device 10 includes an inflatable receptacle 12 deployable between a closed insertion state (Figure 1) and an open state (Figure 2), the open state being configured for substantially enveloping the ventricular portion of the heart in a manner shown in Figure 5. A number of support elements 14 are mechanically linked to inflatable receptacle 12 so as to  
10      10      deploy the inflatable receptacle to its open state. Support elements 14 are preferably implemented as flexible rods resiliently biased to return to a pre-defined shape. These rods serve both to push forward the inflatable receptacle during introduction of the device into a patient and to open it into a correctly positioned and shaped configuration to substantially envelope the ventricular portion of the  
15      15      heart within the pericardium.

In a preferred implementation, the configuration of rods 14 is chosen to provide unguided deployment around the heart in a manner to be described below. This allows "blind" insertion of the device by a minimal invasive procedure. The term "blind" is used in this context to refer to a minimal invasive procedure  
20      20      performed without simultaneous use of imaging techniques for positioning. The minimal invasive nature of the deployment makes the use of the device practical even with patients not sufficiently fit to undergo major surgery and minimizes the trauma to the patient. The possibility of blind insertion renders the device rapidly

deployable so that it can be used when cardiac assist must be rapidly begun or in emergency procedures such as CPR.

It should be appreciated that the device of the present invention may be used in a wide range of circumstances where cardiac assist is required. The device 5 generally serves a "bridge to recovery" or "bridge to surgery" role for periods which may range from a matter of minutes or hours up to weeks or even several months.

In keeping with the normal definitions of "minimal invasive" procedures, the insertion state of device 10 preferably exhibits a maximum diameter of no more 10 than about 4 cm, and typically between about 1.5 and about 2.5 cm. This ensures minimum possible trauma to the patient as a result of the procedure.

Turning now to the features of device 10 in more detail, inflatable receptacle 12 includes at least one, and typically 4-8, inflatable chambers 16 extending along the length of receptacle 12. A preferred arrangement of these 15 chambers may be seen in Figure 3. The inner wall 18 of each chamber is elastically stretchable while the outer wall 20 is relatively non-stretchable. These properties may be achieved either by employing different materials for the inner and outer walls or by making outer wall 20 relatively thick compared to inner wall 18. In either case, the materials may readily be selected from a range of polymer 20 materials conventionally used for surgical devices. The result of these properties is that, when chambers 16 are inflated, the primary effect is directed inward so as to exert pressure on a part of the heart located within inflatable receptacle 12.

It should be noted that chambers 16 may be commonly or independently controlled. By way of example, in certain applications, assistance may be desired exclusively, or primarily, for the left ventricle to prevent excess pressure to the lungs. For such applications, a left/right subdivision of the fluid supply to 5 chambers 16 is required.

As already mentioned, support elements 14 are preferably implemented as flexible rods which exhibit a "memory" for a predefined shape. Usually, at least three rods, and typically between 4 and 8, are used to achieve the desired opening action. The rods may be of metal or polymer material, or a combination thereof, 10 and may have any cross-sectional shape.

A number of possibilities exist for the positioning of rods 14. They may be included within either inner wall 18 or outer wall 20, or may form part of the interconnection between adjacent chambers 16 as shown here. A further possibility is positioning of rods 14 within the volume contained by chambers 16. In all of 15 these cases, especially where rods 14 are made from metal, they are preferably surrounded by a layer of polymer material to avoid direct contact between the rods and body tissue.

A further feature of certain preferred embodiments of device 10 is the provision of a retractable suction clamping system 44. Suction clamping system 44 20 is used to clamp device 10 temporarily in position relative to the heart during deployment, then being released and retracted away from the heart once correct positioning of receptacle 12 around the heart has been achieved. As may best be seen by referring ahead to Figure 7, suction clamping system 44 includes a suction

cup 46 mounted on the end of a tube 48 positioned axially within device 10 in its storage state. Tube 48 is provided with both a suction connection and a mechanical connection for displacing suction clamping system 44 axially. The operation of suction clamping system 44 will be described below with reference to Figures 9A-  
5 9D.

It is a particular feature of most preferred implementations of the present invention that device 10 features a retaining mechanism 22 operable during a minimal invasive procedure to retain inflatable receptacle 12 in a position enveloping the ventricular portion of the heart. Retaining mechanism 22 is  
10 configured to be deployed around a supraventricular portion of the heart, i.e., a portion of the heart lying above the ventricular portion, where the reduced diameter of the heart allows secure retention as shown in Figure 5. At the same time, as mentioned above, it is important that inflatable receptacle 12 does not exert pressure on the atria. To this end, retaining mechanism 22 is preferably  
15 mechanically linked to inflatable receptacle 12 via a non-inflatable web 24. The word "web" is used herein to refer to any continuous or discontinuous flexible layer which forms a mechanical link between retaining mechanism 22 and inflatable receptacle 12. Typically, web 24 is implemented as a single layer continuation of the outer walls of chambers 16. Details of two possible  
20 implementations of retaining mechanism 22 will be described below with reference to Figures 10 and 11.

Insertion and subsequent control of device 10 is controlled through a base connector 26. Base connector 26 houses fluid supply conduits 28, as seen in Figure

4, which connect to the root of each inflatable chamber 16. These conduits preferably have relatively large areas to minimize resistance for a pulsating fluid supply. Base connector 26 additionally provides a mechanical connection to rods 14 and houses retaining mechanism connections 30.

5       Externally, base connector 26 preferably also has guide features such as guide slots 32 for guiding the base connector in a straight path within a trocar tube to be described below, and a projecting handle 34 which allows manual displacement of device 10 along the trocar and provides one or more external fluid connection port 36 as well as a retaining mechanism control 38.

10      Deployment of device 10 is preferably performed as a minimal invasive procedure by use of a trocar. A preferred trocar design and other details of the deployment procedure will now be described with reference to Figures 6-9:

Turning first to Figures 6 and 7, there is shown a two stage trocar made up of an insertion stage 40 for insertion through a small incision and a storage stage 15 42. Second stage 42 has a substantially cylindrical form within which device 10 may be retracted, thereby confining device 10 to an initial closed insertion state as shown in Figure 1. Handle 34 of base connector 26 projects through a longitudinal slot 44 in second stage 42 to provide a manually operable slide for displacing device 10 along the length of the trocar. At the same time, handle 34 provides 20 accessible connections for the various fluid, suction and mechanical connections employed to deploy and operate device 10. Storage stage 42 is preferably attached to insertion stage 40 through a clip-on connection 50 so as to be removable after deployment and replaceable for withdrawal of the device.

Insertion stage 40 preferably includes a curved hollow guide portion 52 for insertion within the patient. Guide portion 52 is preferably formed with a turn angle of between about 15° and about 30°, and typically about 20°. In other words, guide portion 52 redirects device 10 during deployment by an angle of about 20°  
5 from the extensional direction of storage stage 42. This provides the correct alignment for deployment of the device curving under the sternum to an initial position facing the apex of the heart. A shield 54 is preferably deployed around insertion stage 40. This serves to delimit the fully inserted position of the trocar, as well as providing a seal around the incision.

10 Figure 8 shows the deployed state of the device within insertion stage 40 after removal of storage stage 52. In this state, the part of the device projecting from the body is small and can be secured by various techniques such as one or more belt around the patient.

The procedure for deploying device 10 will now be described with reference  
15 to Figures 9A-9D. Initially, insertions stage 40 is introduced through an incision in the outer wall of the abdomen just below the sternum 55 and through a corresponding incision through the lower part of the pericardium 56 in the region near the diaphragm. At this stage, tube 48 is displaced forward and has suction applied so that suction cup 46 clamps to the heart 58 at or near its apex. This  
20 clamping effect maintains the spacial relation between the insertion stage 40 of the trocar and the heart during the rest of the deployment procedure.

Then, handle 34 is gently eased forward along slot 44, thereby advancing device 10 out through guide portion 52. As support elements 14 emerge, their

resilient memory urges them to their predefined diverging form, thereby opening inflatable receptacle 12 to its open form. The simultaneous advancing and opening action brings the device through the stages shown in Figures 9B and 9C until it substantially envelopes the ventricular portion of the heart. At the same time, the 5 distal part of retaining mechanism 22 and its associated non-inflatable web 24 extend over part of the superventricular portion of the heart, primarily on the dorsal side which is not obstructed by blood vessels.

Once the device reaches its fully deployed state, retaining mechanism 22 is tightened around the superventricular portion as shown in Figure 9D. This is 10 sufficient to firmly retain the heart within inflatable receptacle 12 without unduly restricting the normal movement of the heart. The suction clamping is then released and tube 48 withdrawn so that suction cup 46 has no effect on the subsequent operation of the device.

It will be clear that removal of the device is achieved by a simple reversal of 15 the steps described, except that the suction clamping feature is superfluous during removal. The resilient nature of support elements 14 provides an inherent self-opening feature which facilitates opening of retaining mechanism 22.

Turning now briefly to Figure 10, this shows a first preferred implementation of retaining mechanism 22. In this case, the tightening action is 20 achieved by a drawstring mechanism in which a draw-string or wire 60 extends along a pair of sleeves 62 which extend up to near the opening of receptacle 12. In this case retaining mechanism control 38 may be implemented as a simple manual

or automated drawstring connection which pulls on the wire 60, thereby shortening the circumference of the opening between the ends of sleeves 62.

An alternative implementation of retaining mechanism 22 is shown in Figure 11. Here an inflatable collar 64 may be inflated through an independent 5 fluid supply conduit 66 to lodge against the supraventricular portion of the heart.

Turning now briefly to the operation of deployed device 10, the device is driven by a pulsed source of fluid, typically in the form of a pulsatile pump. In an implementation for very short term emergency use, a manual hand pump may be used. The operating fluid may be liquid, gas or a mixture thereof, and is preferably 10 sterile.

Timing of the pulses is preferably performed according to one of three modes. Firstly, in a CPR case where unaided heart activity has stopped, pulses are generated either manually or according to suitable pulse timing circuitry of any kind. Secondly, where either a natural or induced heart beat is detectable, the 15 pulses are preferably synchronized with the heart, either through an electrocardiogram (ECG) signal or by direct synchronization with a pacemaker or defibrillator. Details of how to achieve ECG synchronization may be found in a number of sources such as the aforementioned U.S. Patent No. 5,713,954 to Rosenberg et al. which is hereby incorporated by reference in its entirety. In a third 20 mode, the basic synchronization with an ECG signal is enhanced by closed loop feedback. Suitable sources for feedback to provide a measure of the effectiveness of the cardiac assist include, but are not limited to, measurements of the cardiac output, the ejection fraction, or the systolic blood pressure. These quantities may

be measured by any conventional techniques. Clearly, to implement these various modes, suitable hardware or hardware/software combinations are provided. These typically include a microprocessor unit operating appropriate software under a suitable operating system as is well known in the art.

5        Finally, it should be appreciated that the device of the present invention may be used to advantage in conjunction with a range of other devices and techniques. Examples include, but are not limited to, use in combination with a pacemaker, defribulator, ventilator systems or CPR systems. For such applications, one or more electrodes 70 may be incorporated into the inner surface for inflatable  
10      receptacle as shown schematically in Figure 3.

It will be appreciated that the above descriptions are intended only to serve as examples, and that many other embodiments are possible within the spirit and the scope of the present invention.

## WHAT IS CLAIMED IS:

1. A cardiac massage device deployable by a minimal invasive procedure for applying pulsatory pressure to a ventricular wall of a heart, the device comprising:
  - (a) an inflatable receptacle deployable between a closed insertion state and an open state, said open state being configured for substantially enveloping the ventricular portion of the heart; and
  - (b) a plurality of support elements mechanically linked to said inflatable receptacle so as to deploy said inflatable receptacle to said open state.
2. The device of claim 1, wherein said support elements are implemented as flexible rods resiliently biased to return to a pre-defined shape.
3. The device of claim 2, further comprising an elongated hollow structure for storage and deployment of the device, said inflatable receptacle and said flexible rods being retractable within said elongated hollow structure so as to confine said flexible rods to an initial state corresponding to said closed insertion state of said inflatable receptacle, said flexible rods returning to said pre-defined shape as they advance out from said elongated hollow structure.
4. The device of claim 3, wherein said elongated hollow structure forms a storage stage of a two-stage trocar, an insertion stage of said trocar being

configured for insertion through the skin of a patient as part of a minimal invasive surgical procedure.

5. The device of claim 4, wherein said insertion stage of said trocar includes a curved hollow guide portion for insertion within the patient, said curved hollow guide portion providing a turn angle of between about 15° and about 30°.

6. The device of claim 3, further comprising a manually operable mechanism for displacing said inflatable receptacle and said flexible rods along a length of said trocar.

7. The device of claim 1, further comprising a retaining mechanism associated with said inflatable receptacle and configured for deploying around a supraventricular portion of the heart so as to retain said inflatable receptacle in a position enveloping the ventricular portion of the heart.

8. The device of claim 7, wherein said retaining mechanism includes at least one drawstring mechanism.

9. The device of claim 8, wherein said at least one drawstring mechanism is connected to said inflatable receptacle via a non-inflatable web.

10. The device of claim 7, wherein said retaining mechanism includes at least one inflatable retaining element, said inflatable retaining element and said inflatable receptacle being provided with independent fluid supply conduits.

11. The device of claim 1, further comprising a retractable suction clamping system including:

- (a) a suction cup deployed for suction clamping to the apex of the heart; and
- (b) an axially displaceable tube forming a fluid connection with said suction cup,

such that, when said tube is displaced forward and suction is applied to said tube, said suction cup clamps to the heart and, when the suction is released and said tube is displaced rearwards, said suction cup is withdrawn from contact with the heart.

12. The device of claim 1, further comprising at least one electrode deployed on an inner surface of said inflatable receptacle so as to contact the heart when the device is deployed.

13. A cardiac massage device deployable by a minimal invasive procedure for applying pulsatory pressure to a ventricular wall of a heart, the device comprising:

- (a) an inflatable receptacle for substantially enveloping the ventricular portion of the heart, said inflatable receptacle being configured to

exert pulsating pressure against the ventricular wall in response to a pulsating fluid supply; and

- (b) a retaining mechanism associated with said inflatable receptacle and configured for deploying around a supraventricular portion of the heart so as to retain said inflatable receptacle in a position substantially enveloping the ventricular portion of the heart.

14. The device of claim 13, wherein said retaining mechanism includes at least one drawstring mechanism.

15. The device of claim 14, wherein said at least one drawstring mechanism is connected to said inflatable receptacle via a non-inflatable web.

16. The device of claim 13, wherein said retaining mechanism includes at least one inflatable retaining element, said inflatable retaining element and said inflatable receptacle being provided with independent fluid supply conduits.

17. A method of minimal invasive deployment of a cardiac massage device around the ventricular portion of the heart of a patient, the method comprising the steps of:

- (a) introducing the device in a compact state through the pericardium in the region near the diaphragm; and
- (b) opening the device to form a receptacle which substantially envelopes the ventricular portion of the heart.

18. The method of claim 17, wherein said opening of the device occurs through a plurality of resiliently biased flexible rods deployed within the device returning to a pre-defined shape.

19. The method of claim 17, further comprising application of suction clamping to the surface of the heart during deployment of the device, and removal of the suction clamping once the device is properly deployed.

20. The method of claim 17, further comprising deploying a retaining mechanism around a supraventricular portion of the heart so as to retain said receptacle in a position enveloping the ventricular portion of the heart.

21. The method of claim 17, wherein said compact state exhibits a maximum diameter of no more than about 4 cm.

22. The method of claim 17, wherein said compact state exhibits a maximum diameter of between about 1.5 and about 2.5 cm.

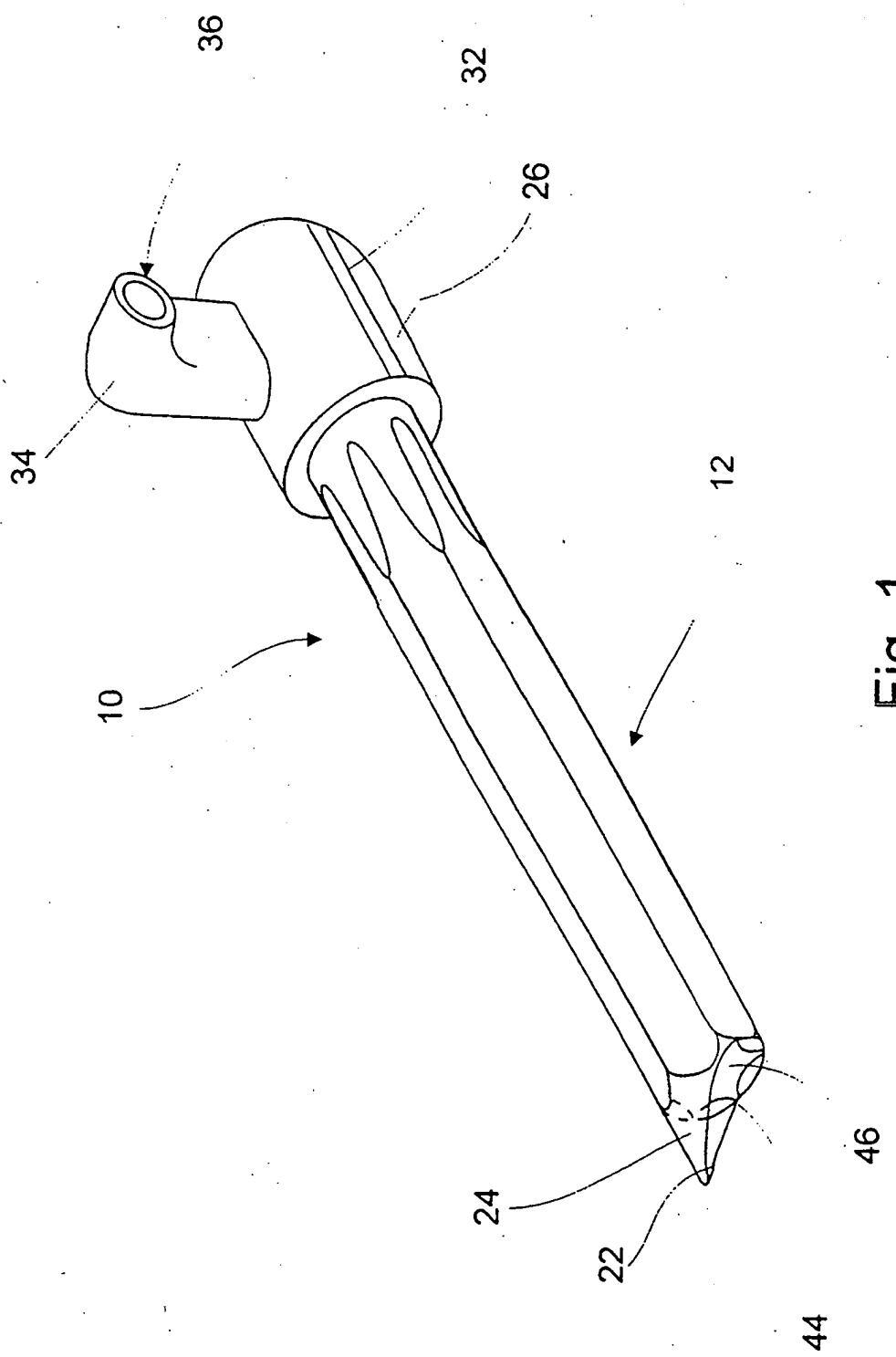


Fig. 1

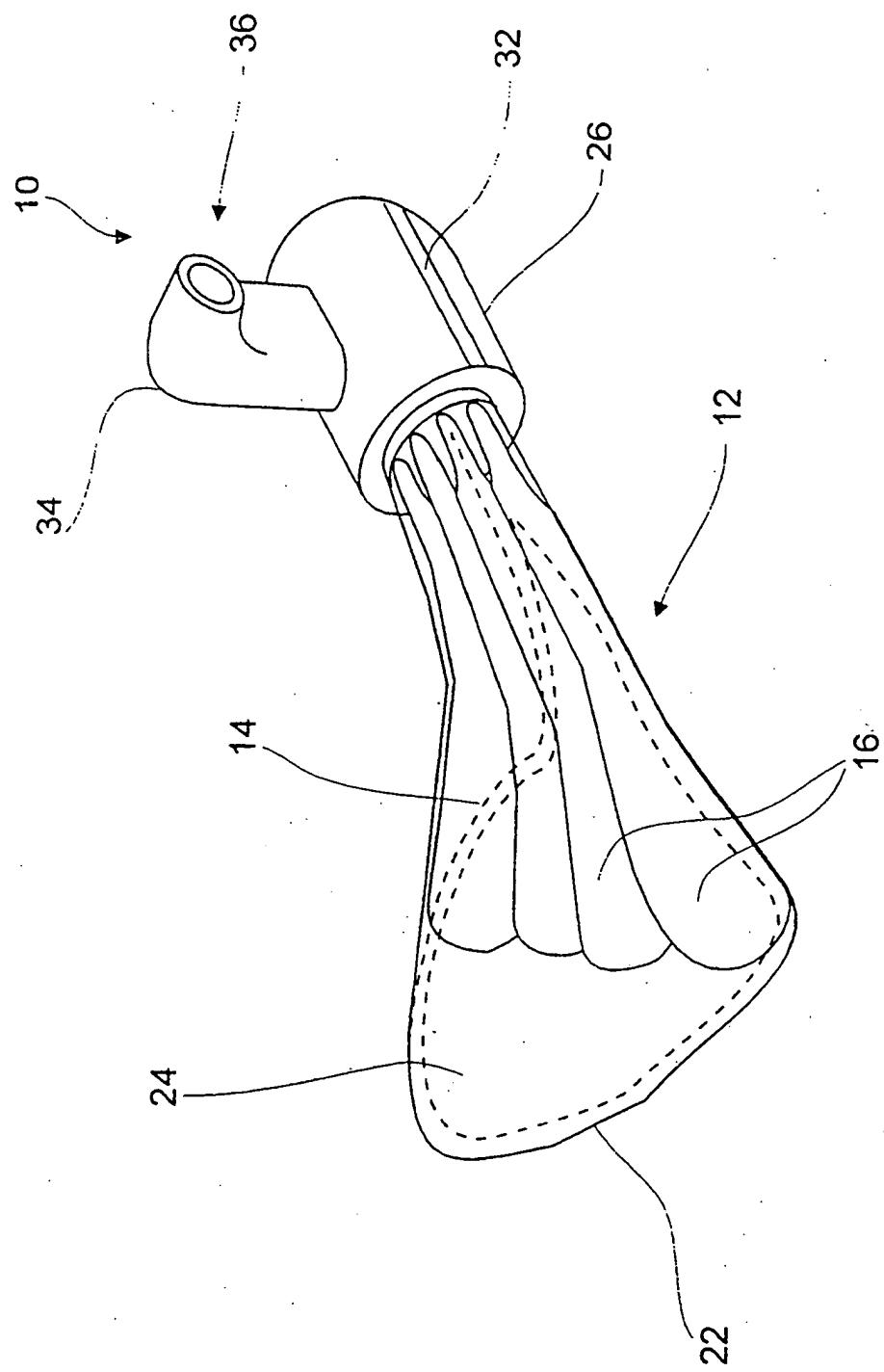


Fig. 2

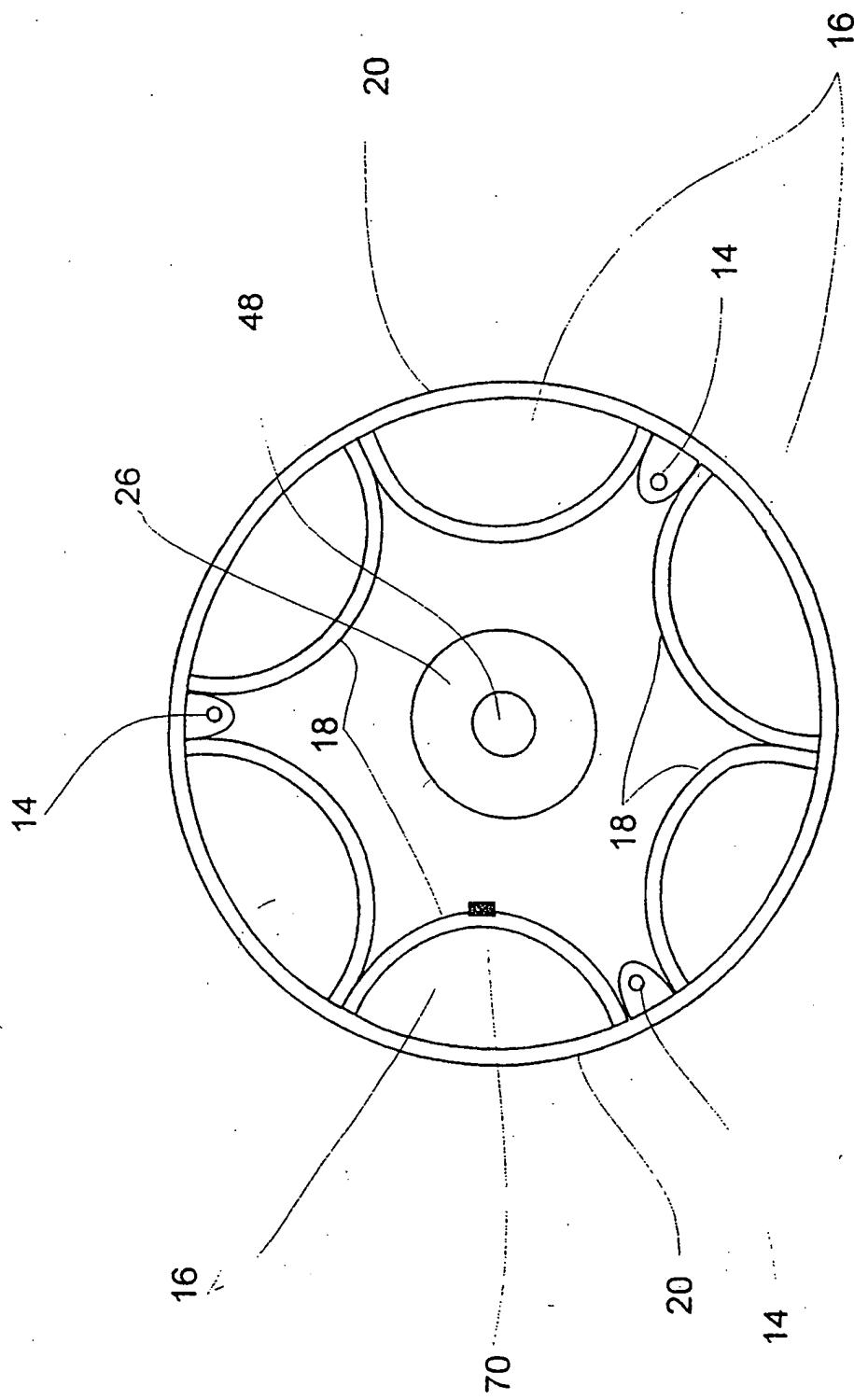


Fig. 3

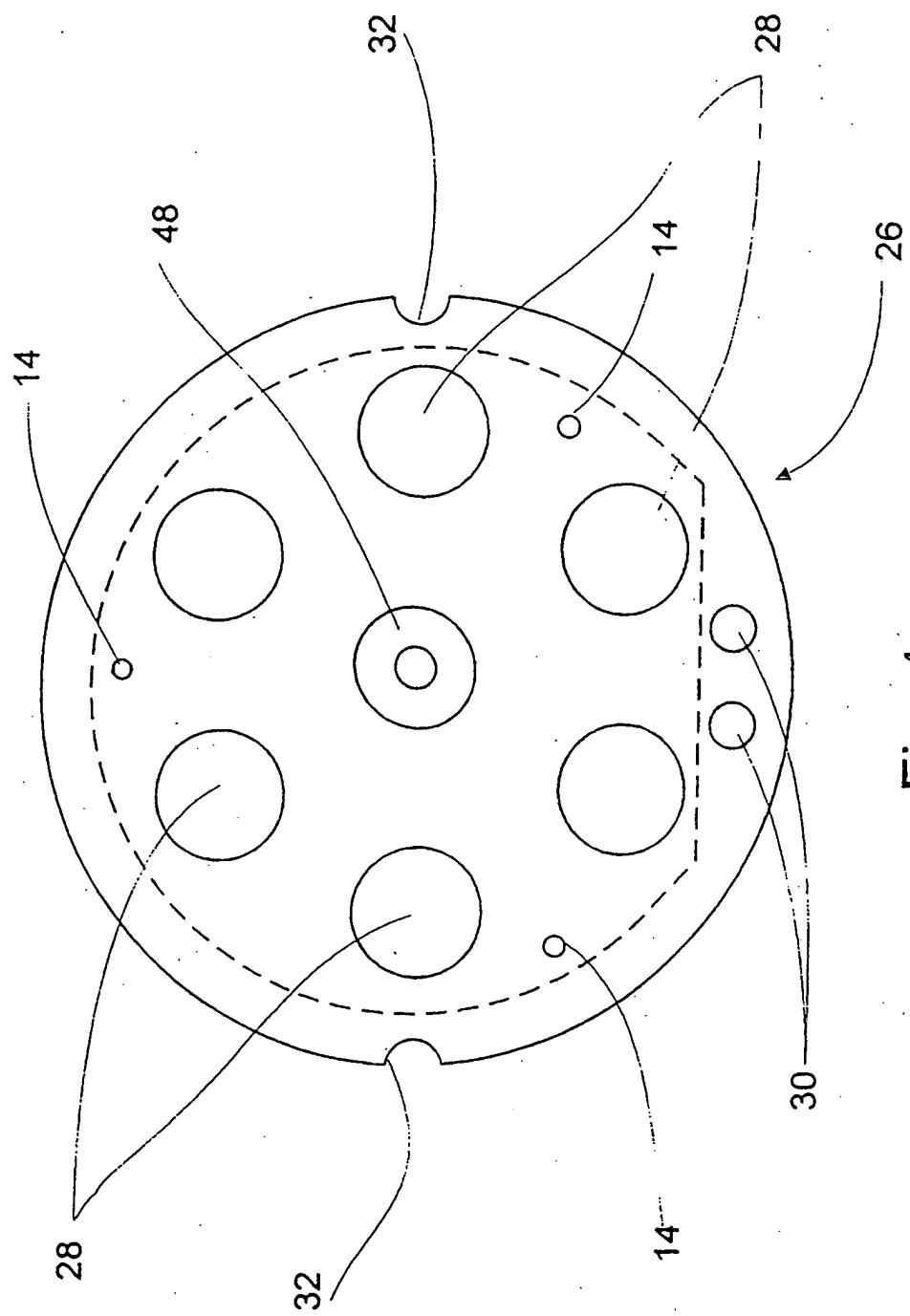


Fig. 4

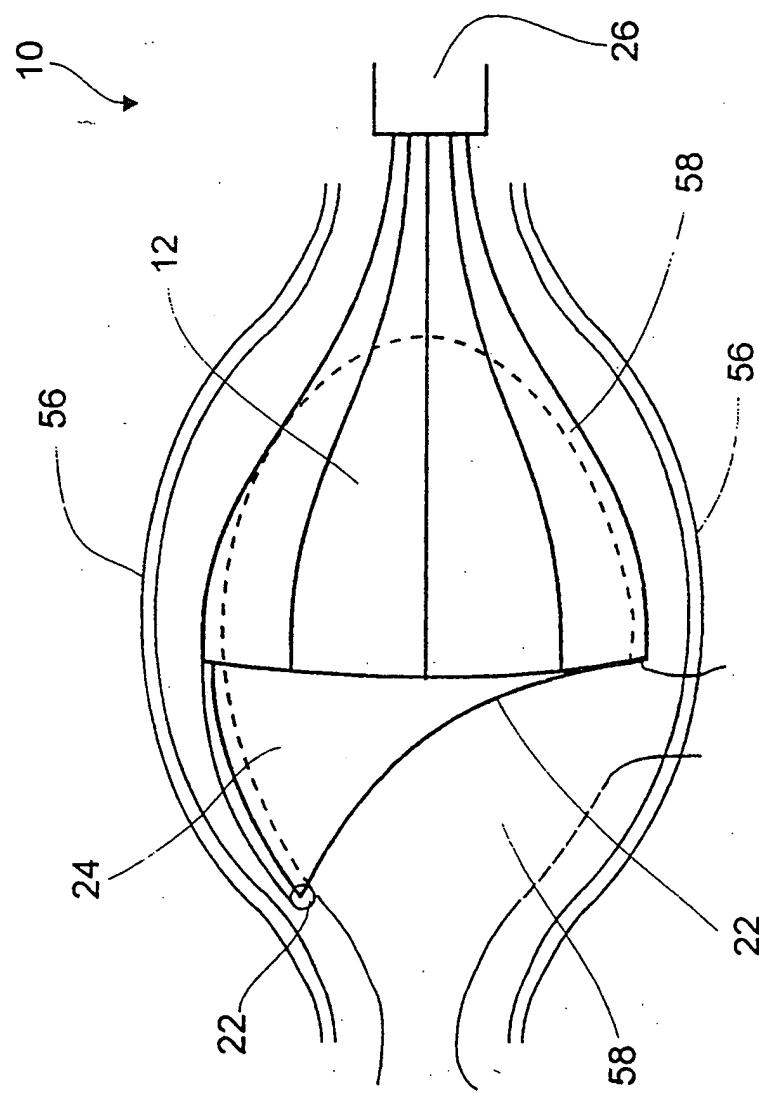


Fig. 5

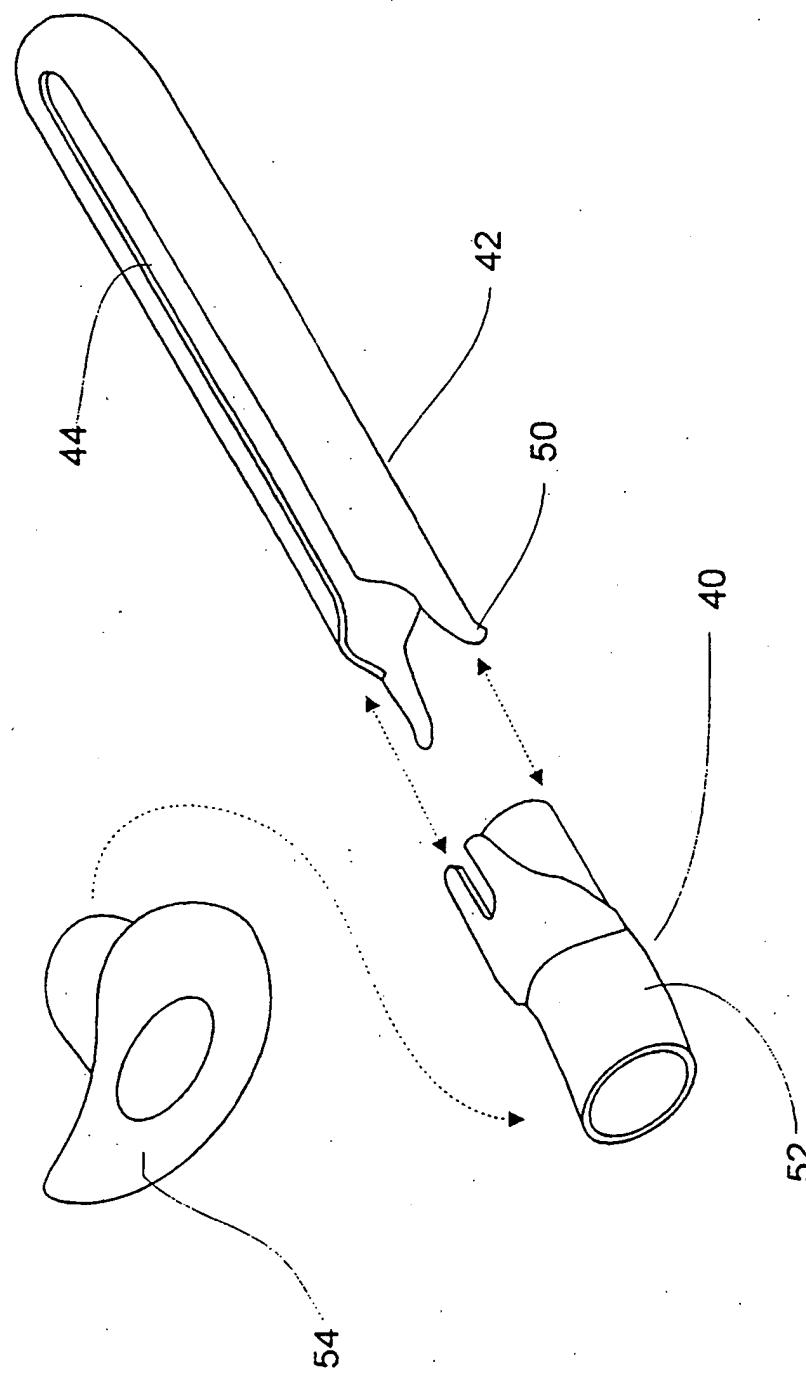


Fig. 6

7/11

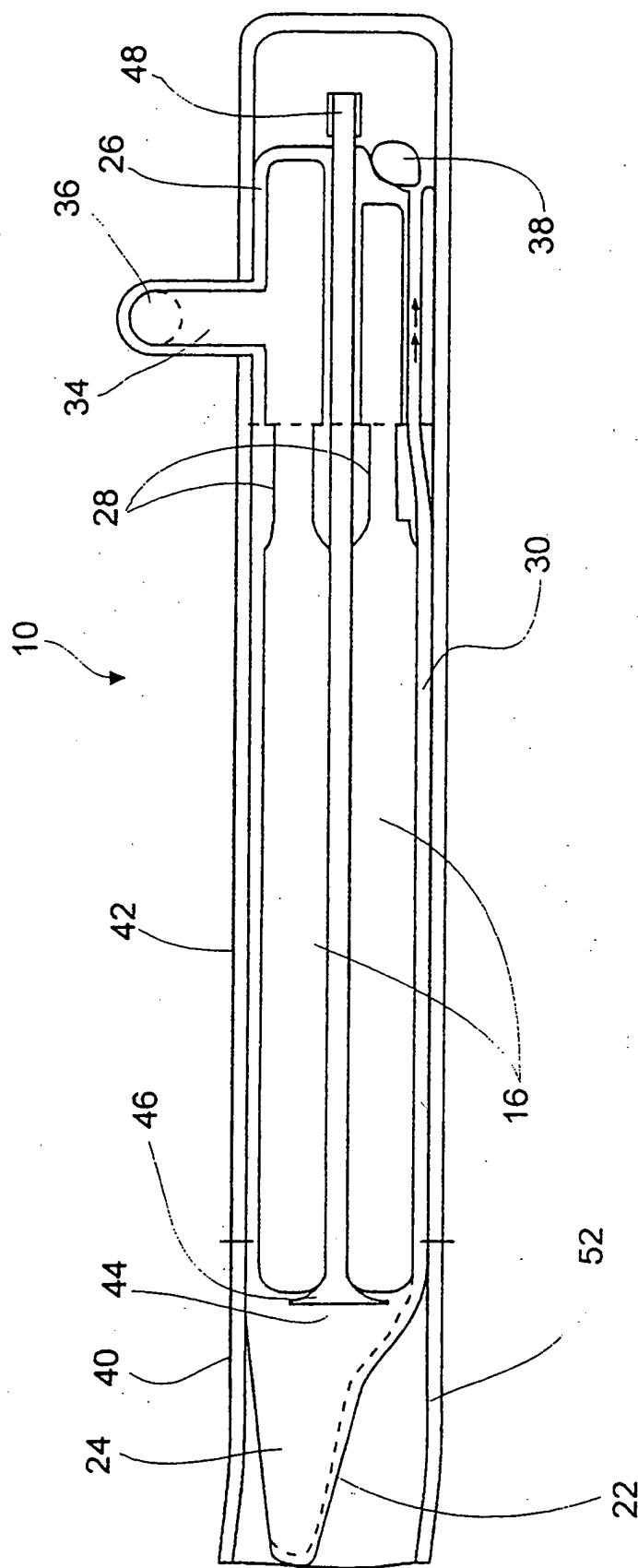


Fig. 7

8/11

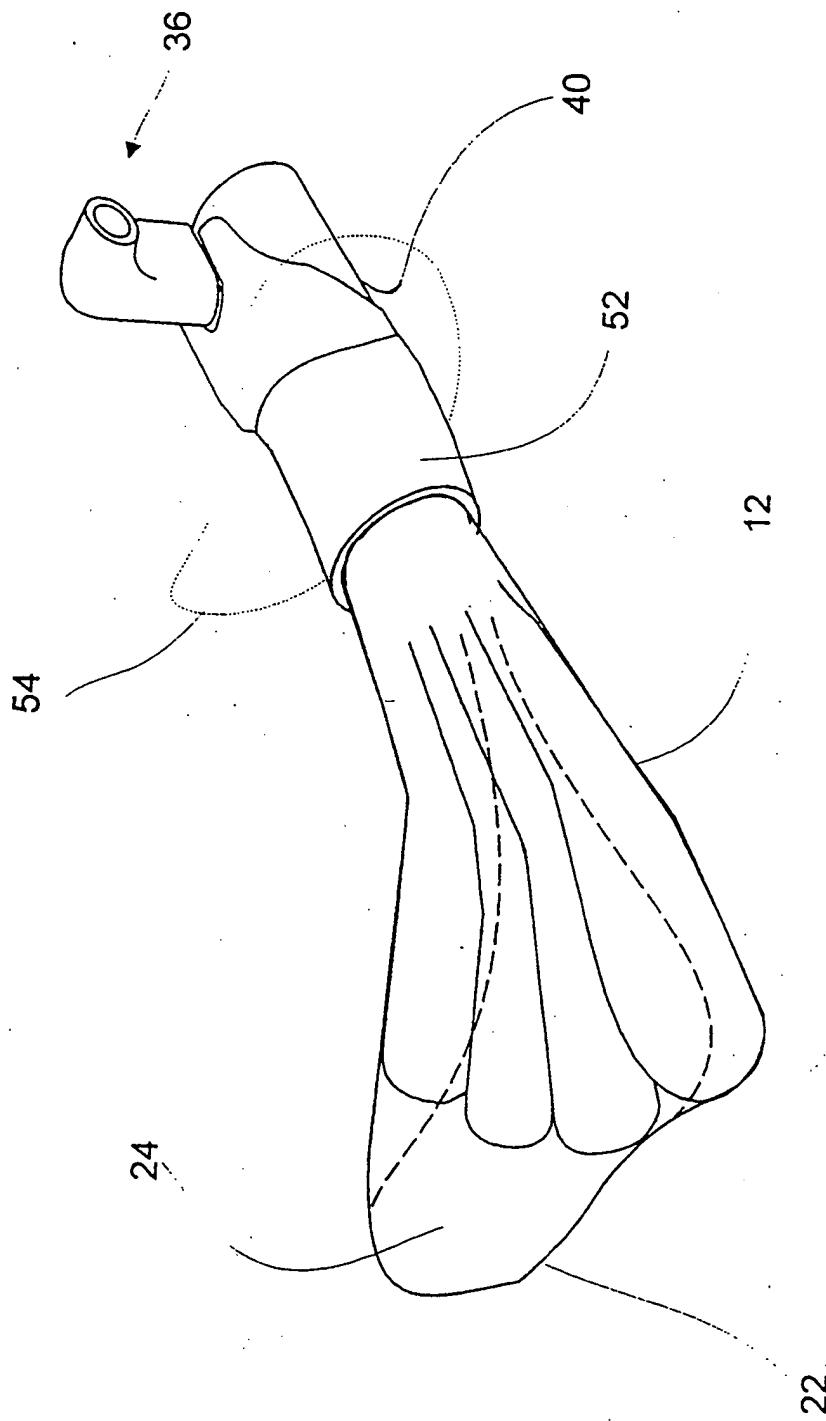


Fig. 8

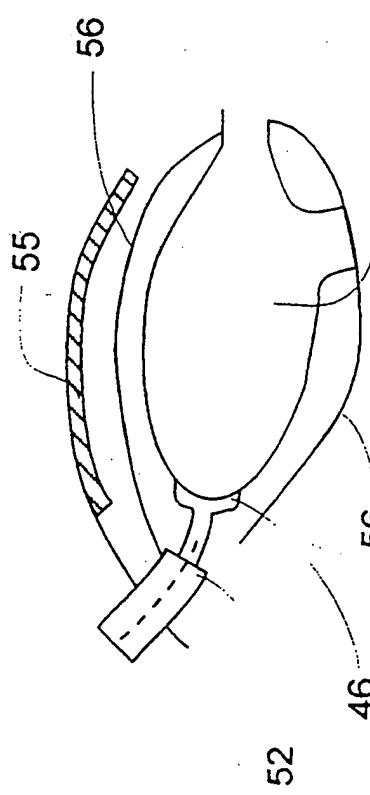


Fig. 9a

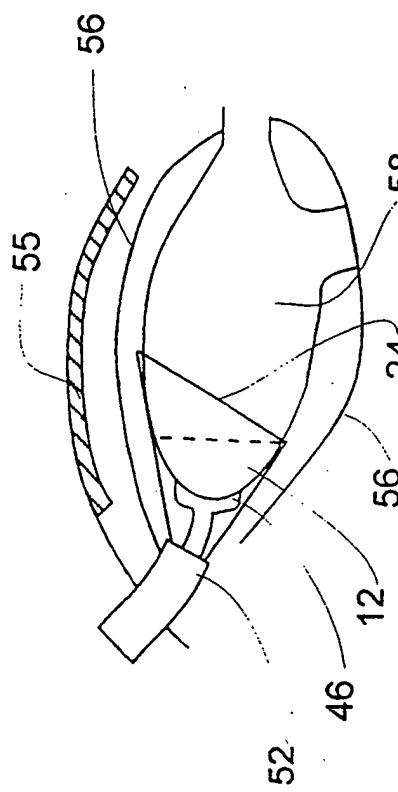


Fig. 9b

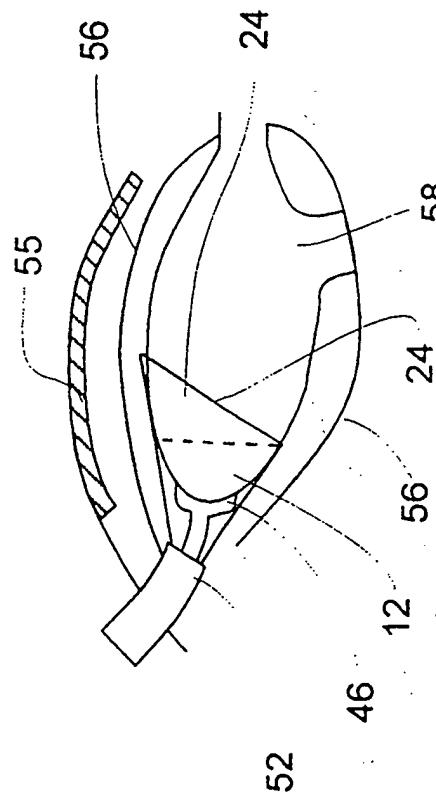


Fig. 9c

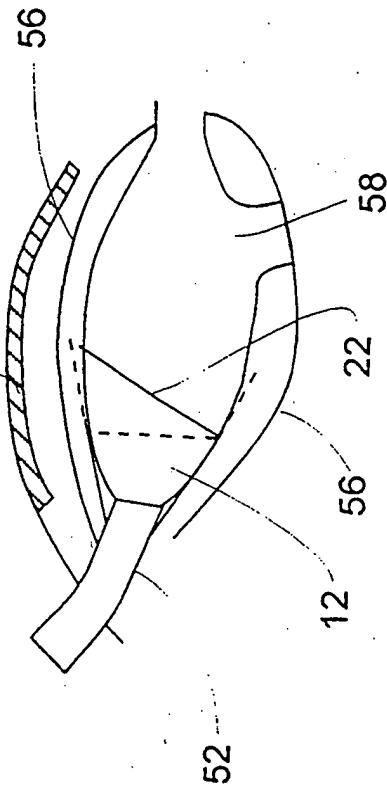


Fig. 9d

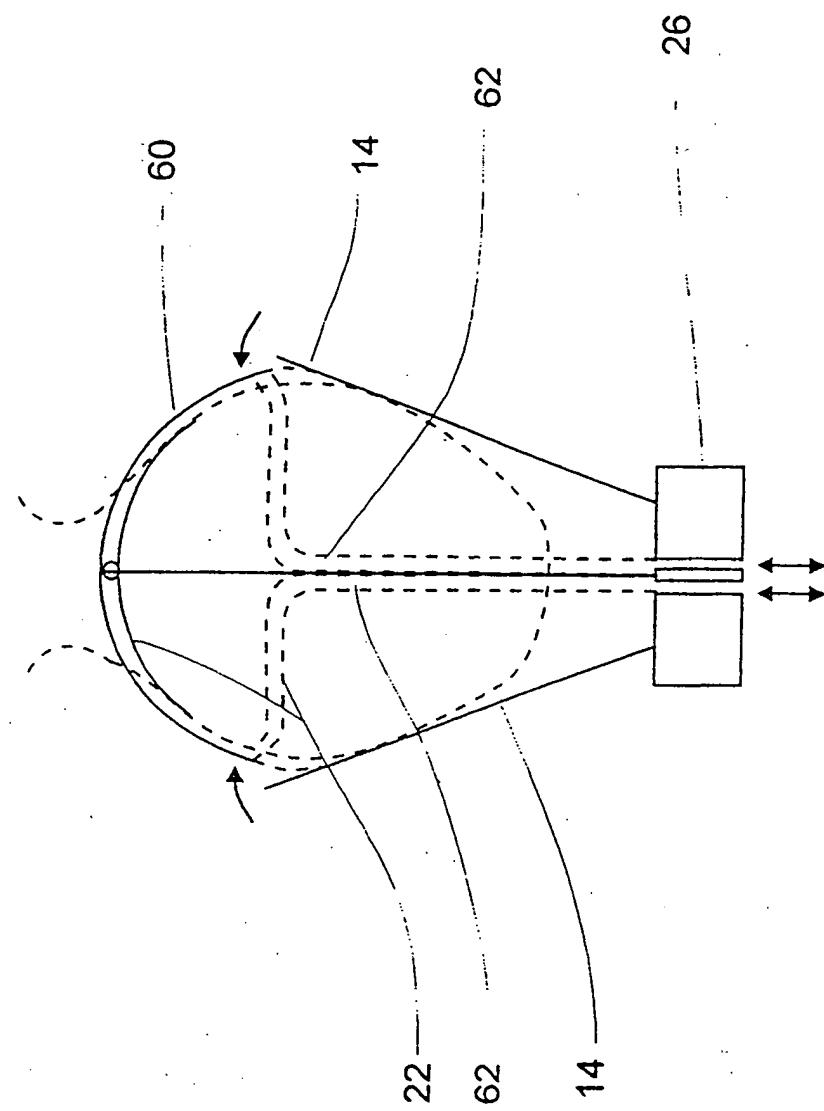


Fig. 10

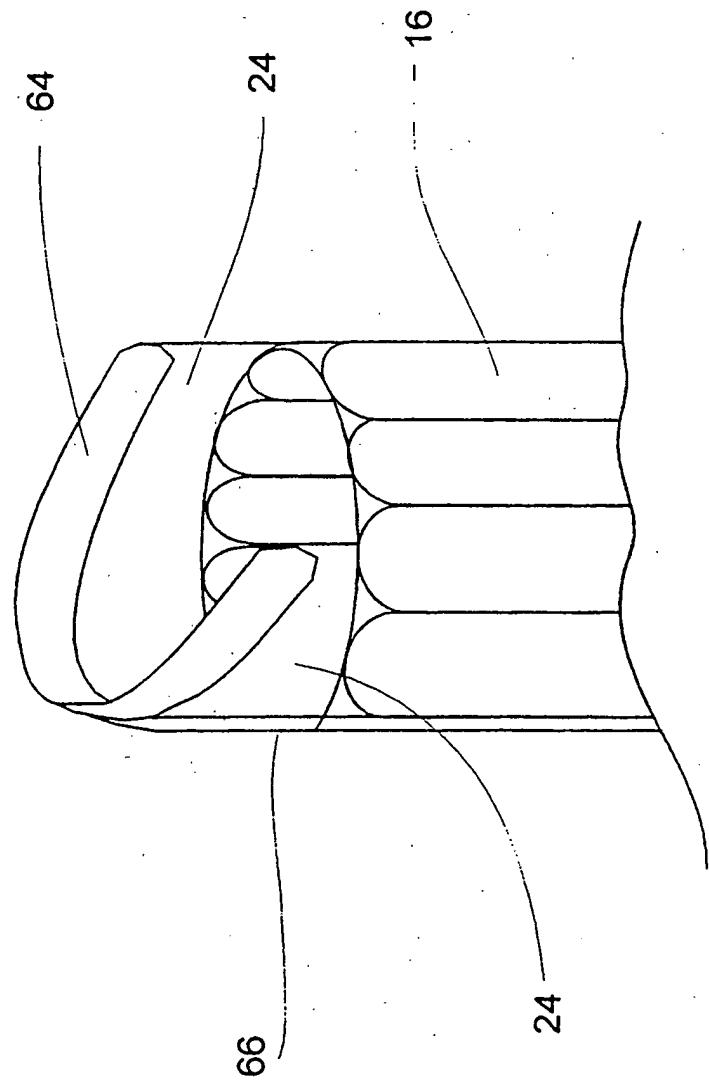


Fig. 11

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/IL98/00318

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61B 17/00; A61F 2/04  
US CL : 600/17; 623/3

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 600/17; 623/3

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5,707,336 A (RUBIN) 13 January 1998 (13.01.98), entire patent.	1-22

Further documents are listed in the continuation of Box C.

See patent family annex.

Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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*O* document referring to an oral disclosure, use, exhibition or other means		
*P* document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

10 NOVEMBER 1998

Date of mailing of the international search report

16 DEC 1998

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